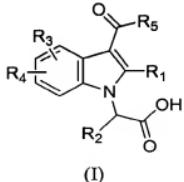


This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Currently amended) Compounds of formula (I):



wherein:

R₁ is hydrogen, C₁-C₆ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, or C₁-C₃ perfluoroalkyl, wherein the alkyl and cycloalkyl groups may be optionally substituted with halogen, -CN, C₁-C₆ alkoxy, -OH, -NH₂, or -NO₂;

R₂ is hydrogen, C₁-C₈ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, thiienyl, -CH₂-thienyl, furanyl, CH₂-furanyl, oxazoyl, CH₂-oxazoyl, phenyl, benzyl, CH₂-naphthyl, wherein the alkyl group and the rings of the cycloalkyl, thiienyl, furanyl, oxazoyl, phenyl, benzyl, and naphthyl groups may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₁-C₃ perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, -OCHF₂, -CN, -COOH, -CH₂CO₂H, -C(O)CH₃, -CO₂R₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂;

R₃ is hydrogen, halogen, C₁-C₆ alkyl, C₁-C₃ perfluoroalkyl, C₁-C₆ alkoxy, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, -NH₂, or -NO₂;

R₄ is C₃-C₈ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, thiienyl, furanyl, oxazoyl, phenyl, benzofuran-2-yl, benzo[b]thien-2-yl, benzo[1,3]dioxol-5-yl, naphthyl, wherein the alkyl groups and the rings of the cycloalkyl, thiienyl, furanyl, oxazoyl, phenyl, benzofuranyl, benzothienyl, and naphthyl groups may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₁-C₃

perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, -OCHF₂, -CN, -COOH, CH₂CO₂H, -C(O)CH₃, -C(O)OR₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂;

R₅ is C₁-C₈ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, **pyridinyl**, **CH₂-pyridinyl**, **thienyl**, **CH₂-thienyl**, **furanyl**, **CH₂-furanyl**, **oxazoyl**, **CH₂-oxazoyl**, **phenyl**, **benzyl**, **benzo[b]furan-2-yl**, **benzo[b]thien-2-yl**, **benzo[1,3]dioxol-5-yl**, **naphthyl**, **CH₂-naphthyl**, **9H-fluoren-1-yl**, **9H-fluoren-4-yl**, **9H-fluoren-9-yl**, **9-fluorenone-1-yl**, **9-fluorenone-2-yl**, **9-fluorenone-4-yl**, **CH₂-9H-fluoren-9-yl**, wherein the alkyl group and the rings of the cycloalkyl, **pyridinyl**, **thienyl**, **furanyl**, **oxazoyl**, **phenyl**, **benzyl**, **benzofuranyl**, **benzothienyl**, **naphthyl**, **fluorenlyl**, and **fluorenone groups** may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₃-C₆ cycloalkyl, C₁-C₃ perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, phenoxy, -OCHF₂, -CN, -COOH, -CH₂CO₂H, -C(O)CH₃, -CO₂R₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂, wherein the phenoxy group may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, or C₁-C₃ perfluoroalkyl; and

R₆ is C₁-C₆ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, or benzyl; or a pharmaceutically acceptable salt or ester form thereof.

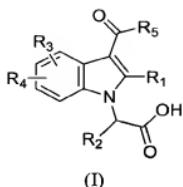
2. (Currently amended) The compound of claim 1 wherein R₁-R₃ and R₅-R₆ are as defined in claim 1, and R₄ is **thienyl**, **furanyl**, **oxazoyl**, **phenyl**, **benzo[b]furan-2-yl**, **benzo[b]thien-2-yl**, **benzo[1,3]dioxol-5-yl**, or **naphthyl**, wherein the rings of the **thienyl**, **furanyl**, **oxazoyl**, **phenyl**, **benzofuranyl**, **benzothienyl**, and **naphthyl** groups may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₁-C₃ perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, -OCHF₂, -CN, -COOH, -CH₂CO₂H, -C(O)CH₃, -C(O)OR₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂.

3. (Original) The compound of claim 1 which is [3-(4-chlorobenzoyl)-5-(4-chlorophenyl)-1*H*-indol-1-yl]acetic acid, or a pharmaceutically acceptable salt or ester form thereof.

4. (Cancelled)

5. (Original) The compound of claim 1 which is [3-(4-chlorobenzoyl)-5-(4-methylphenyl)-1*H*-indol-1-yl]-acetic acid, or a pharmaceutically acceptable salt or ester form thereof.

6. (Currently amended) A method of inhibiting in a mammal plasminogen activator inhibitor type I, comprising administering to a mammal in need thereof a therapeutically effective amount of a compound of formula (I)



wherein:

R₁ is hydrogen, C₂-C₆ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, or C₁-C₃ perfluoroalkyl, wherein the alkyl and cycloalkyl groups may be optionally substituted with halogen, -CN, C₁-C₆ alkoxy, -OH, -NH₂, or -NO₂;

R₂ is hydrogen, C₁-C₈ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, thiienyl, -CH₂-thienyl, furanyl, -CH₂-furanyl, oxazoyl, CH₂-oxazoyl, phenyl, benzyl, CH₂-naphthyl, wherein the alkyl group and the rings of the cycloalkyl, thiienyl, furanyl, oxazoyl, phenyl, benzyl, and naphthyl groups may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₁-C₃ perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, -OCHF₂, -CN, -COOH, -CH₂CO₂H, -C(O)CH₃, -CO₂R₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂;

R₃ is hydrogen, halogen, C₁-C₆ alkyl, C₁-C₃ perfluoroalkyl, C₁-C₆ alkoxy, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, -NH₂, or -NO₂;

R₄ is C₂-C₈ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, thiényl, furanyl, oxazoyl, phenyl, benzo[b]furan-2-yl, benzo[b]thien-2-yl, benzo[1,3]dioxol-5-yl, naphthyl, wherein the alkyl groups and the rings of the cycloalkyl, thiényl, furanyl, oxazoyl, phenyl, benzofuranyl, benzothienyl, and naphthyl groups may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₁-C₃ perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, -OCHF₂, -CN, -COOH, CH₂CO₂H, -C(O)CH₃, -C(O)OR₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂;

R₅ is C₁-C₈ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, pyridinyl, -CH₂-pyridinyl, thiényl, CH₂-thienyl, furanyl, CH₂-furanyl, oxazoyl, CH₂-oxazoyl, phenyl, benzyl, benzo[b]furan-2-yl, benzo[b]thien-2-yl, benzo[1,3]dioxol-5-yl, naphthyl, CH₂-naphthyl, 9H-fluoren-1-yl, 9H-fluoren-4-yl, 9H-fluoren-9-yl, 9-fluorenone-1-yl, 9-fluorenone-2-yl, 9-fluorenone-4-yl, CH₂-9H-fluoren-9-yl, wherein the alkyl group and the rings of the cycloalkyl, pyridinyl, thiényl, furanyl, oxazoyl, phenyl, benzyl, benzofuranyl, benzothienyl, naphthyl, fluorenyl, and fluorenone groups may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, C₃-C₆ cycloalkyl, C₁-C₃ perfluoroalkyl, -O-C₁-C₃ perfluoroalkyl, -S-C₁-C₃ perfluoroalkyl, C₁-C₃ alkoxy, phenoxy, -OCHF₂, -CN, -COOH, -CH₂CO₂H, -C(O)CH₃, -CO₂R₆, -C(O)NH₂, -S(O)₂CH₃, -OH, -NH₂, or -NO₂, wherein the phenoxy group may be optionally substituted by from 1 to 3 groups selected from halogen, C₁-C₃ alkyl, or C₁-C₃ perfluoroalkyl; and

R₆ is C₁-C₆ alkyl, C₃-C₆ cycloalkyl, -CH₂-C₃-C₆ cycloalkyl, or benzyl; or a pharmaceutically acceptable salt or ester form thereof.

7. (Withdrawn) A pharmaceutical composition comprising a compound of claim 1 and a pharmaceutical carrier.

8. (Withdrawn) A method for treatment of thrombosis or fibrinolytic impairment in a mammal, the method comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

9. (Withdrawn) A method of Claim 8 wherein the thrombosis or fibrinolytic impairment is associated with formation of atherosclerotic plaques, venous and arterial thrombosis, myocardial ischemia, atrial fibrillation, deep vein thrombosis, coagulation syndromes, pulmonary fibrosis, cerebral thrombosis, thromboembolic complications of surgery or peripheral arterial occlusion.

10. (Withdrawn) A method for the treatment of peripheral arterial disease in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

11. (Withdrawn) A method for the treatment of stroke associated with or resulting from atrial fibrillation in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

12. (Withdrawn) A method for the treatment of deep vein thrombosis in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

13. (Withdrawn) A method for the treatment of myocardial ischemia in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

14. (Withdrawn) A method for the treatment of a cardiovascular disease caused by noninsulin dependent diabetes mellitus in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

15. (Withdrawn) A method for the treatment of the formation of atherosclerotic plaques in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

16. (Withdrawn) A method for the treatment of chronic obstructive pulmonary disease in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

17. (Withdrawn) A method for the treatment of renal fibrosis in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

18. (Withdrawn) A method for the treatment of polycystic ovary syndrome in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

19. (Withdrawn) A method for the treatment of Alzheimer's disease in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.

20. (Withdrawn) A method for the treatment of cancer in a mammal, comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound of Claim 1.